

ORGANIC ELECTROLYTIC SOLUTION AND LITHIUM BATTERY USING THE SAME

BACKGROUND OF THE INVENTION

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This application claims priority from Korean Patent Application No. 2002-71043, filed on November 15, 2002, in the Korean Intellectual Property Office, the disclosure of which is incorporated herein in its entirety by reference.

10 1. Field of the Invention

The present invention relates to an organic electrolytic solution and a lithium battery containing the same, and more particularly, to an organic electrolytic solution capable of stabilizing lithium metal and improving lithium ionic conductivity, and a lithium battery containing the organic electrolytic solution.

15 2. Description of the Related Art

With the rapid advance of compact portable electronic devices, there is an increasing need for batteries having high energy densities for miniature portable electronic devices.

Lithium sulfur batteries are known as the most promising types of batteries that are capable of satisfying the above requirement over other batteries developed by far due to their high energy density. Lithium and sulfur (S_8) used as active materials in the manufacture of lithium sulfur batteries have an energy density of about 3,830 mAh/g and 1,675 mAh/g, respectively, and are known as being economical and environmentally friendly. However, there has been no successful commercial use of these active materials in battery systems. The reason why it has been difficult to commercialize lithium sulfur batteries lies in the low availability of sulfur as an active material in electrochemical oxidation reactions, which finally leads to low battery capacitance. In addition, the lifespan of batteries can be shortened due to the outflow of sulfur to electrolyte during oxidation and reduction reactions. If an unsuitable electrolytic solution is used, sulfur is reduced and separated as lithium sulfide (Li_2S) that is no longer available in electrochemical reactions.

To resolve these problems, many attempts have been made to optimize the composition of the electrolytic solution. As an example, U.S. Patent No. 6,030,720 discloses use of a mixture of a main solvent such as tetraglyme and a donor solvent

having 15 or greater donor number, such as n,n-diethylacetamide, as an organic solvent of an organic electrolyte.

U.S. Patent No. 5,961,672 discloses use of an organic electrolytic solution of 1 M LiSO_3CF_3 in a mixed solvent of 1,3-dioxolane, diglyme, sulfolane, and diethoxyethane for improved lifespan and safety measures of batteries, wherein a lithium metal anode is coated with a polymeric film.

When a lithium metal electrode is used as an anode of a lithium secondary battery, the lifespan, capacitance, and other properties of the battery degrade compared to using a carbonaceous or graphite electrode. In particular, as a result of repeated charging/discharging cycles, dendrites are separated and grow on the surface of the lithium metal anode, and contact the surface of a cathode, thereby causing shorting out. In addition, the lithium metal corrodes as a result of a reaction with an electrolytic solution at the surface of the lithium anode.

As a solution to these problems, a method of forming a protecting layer on the surface of the lithium metal electrode has been suggested (U.S. Patent Nos. 6,017,651, 6,025,094, and 5,961,672). To be effective, the protecting layer formed on the surface of the lithium electrode should allow lithium ions to pass through itself as well as act as a barrier to prevent an electrolytic solution from contacting the lithium metal of the anode.

In general, this lithium-protecting layer is formed by the reaction of lithium and a protective layer-forming additive contained in the electrolytic solution after the assembly of the battery. However, the protecting layer formed by this method has poor density, so that a considerable amount of electrolytic solution permeates through pores present in the protective layer and undesirably react with lithium metal.

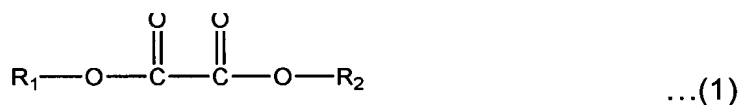
Another method of forming a lithium-protecting layer involves processing the surface of a lithium electrode with nitrogen plasma to form a lithium nitride (Li_3N) layer on the electrode. However, the lithium nitride layer formed by this method includes grain boundaries through which the electrolytic solution easily permeates, is highly likely to decompose when in contact with water, and has a low potential window. Therefore, the lithium nitride layer is impractical to use.

SUMMARY OF THE INVENTION

The present invention provides an organic electrolytic solution that stabilizes, lithium metal due to reduction in reactivity with lithium metal and improves the conductivity of lithium ions.

The present invention also provides a lithium battery providing improved charging/discharging efficiency by using the organic electrolytic solution.

In accordance with an aspect of the present invention, there is provided an organic electrolytic solution comprising: a lithium salt; an organic solvent; and an oxalate compound of formula (1) below:



where R₁ and R₂ are independently selected from hydrogen atom, halogen atom, a hydroxy group, a substituted or unsubstituted C₁-C₂₀ alkyl group, a substituted or unsubstituted C₁-C₂₀ alkoxy group, a substituted or unsubstituted C₁-C₂₀ alkenyl group, a substituted or unsubstituted C₆-C₃₀ aryl group, a substituted or unsubstituted C₆-C₃₀ arylalkyl group, a substituted or unsubstituted C₆-C₃₀ aryloxy group, a substituted or unsubstituted C₂-C₃₀ heteroaryl group, a substituted or unsubstituted C₂-C₃₀ heteroarylalkyl group, a substituted or unsubstituted C₂-C₃₀ heteroaryloxy group, a substituted or unsubstituted C₅-C₂₀ cycloalkyl group, and a substituted or unsubstituted C₂-C₂₀ heterocycloalkyl group.

In accordance with another aspect of the present invention, there is provided a lithium battery comprising: a cathode; an anode; a separator interposed between the cathode and the anode; and the above-described organic electrolytic solution.

BRIEF DESCRIPTION OF THE DRAWINGS

The above and other features and advantages of the present invention will become more apparent by describing in detail exemplary embodiments thereof with reference to the attached drawings in which:

FIGS. 1A and 1B illustrate the interaction between lithium ions and dimethyl oxalate as an example of an oxalate compound of formula (1) above according to the present invention;

FIG. 2 is a graph illustrating change in charging/discharging efficiency with respect to dimethyl oxalate content for lithium batteries manufactured in Examples 1 through 4 according to the present invention and Comparative Example 1;

FIG. 3 is a graph illustrating change in charging/discharging efficiency with respect to diethyl oxalate content for lithium batteries manufactured in Examples 5 through 7 according to the present invention and Comparative Example 1;

FIG. 4 is a graph illustrating change in charging/discharging efficiency with respect to dibutyl oxalate content for lithium batteries manufactured in Examples 8 through 10 according to the present invention and Comparative Example 1;

FIG. 5 is a graph illustrating change in charging/discharging efficiency with respect to bis(4-methylbenzyl) oxalate content for lithium batteries manufactured in Examples 11 through 13 according to the present invention and Comparative Example 1;

FIG. 6 is a graph illustrating change in discharging capacity with respect to the number of charging/discharging cycles for lithium sulfur batteries manufactured in Examples 14, 18, and 19 according to the present invention;

FIG. 7 is a bar graph illustrating charging/discharging efficiency for lithium sulfur batteries manufactured in Examples 14 through 17 according to the present invention and Comparative Examples 1 and 2; and

FIG. 8 is a graph illustrating change in discharging capacity with respect to the number of charging/discharging cycles for lithium sulfur batteries manufactured in Example 14 according to the present invention and Comparative Example 3.

DETAILED DESCRIPTION OF THE INVENTION

Hereinafter, an organic electrolytic solution and a lithium battery using the organic electrolytic solution according to the present invention will be described in detail.

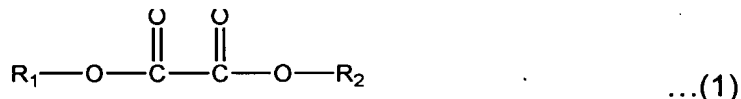
In general, the charging/discharging behaviour of lithium batteries is greatly affected by the properties of a thin film formed on the surface of an electrode. The formation of dendrites on the surface of the lithium anode during charging/discharging cycles causes shorting out of the battery and adversely affects the battery lifespan.

As a lithium battery is charged, a solid electrolyte interface (SEI) is formed on the surface of its anode as a result of decomposition of the electrolytic solution

contained therein. This SEI effectively suppresses side reactions occurring at the anode surface and improves the battery lifespan. However, with repeated charging/discharging cycles of the battery, even the SEI degrades and the electrolytic solution decomposes more and more at the surface of the anode.

Accordingly, in the present invention, an organic electrolytic solution for lithium batteries is prepared by adding an oxalate compound of formula (1) below that is chelatable with lithium ions. The organic electrolytic solution according to the present invention has reduced reactivity to lithium and improved ionic conductivity due to the chelation of lithium ions with the oxalate compound of formula (1) below. Therefore, the organic electrolytic solution according to the present invention can improve charging/discharging efficiency when used in lithium batteries.

When an organic electrolytic solution according to the present invention containing an oxalate compound of formula (1) below is used, especially in a lithium sulfur battery, bonding between lithium ions and sulfide anions is blocked, so that the solubility of sulfide is improved. As a result, a reversible capacity of sulfur is increased.



where R₁ and R₂ are independently selected from hydrogen atom, halogen atom, a hydroxy group, a substituted or unsubstituted C₁-C₂₀ alkyl group, a substituted or unsubstituted C₁-C₂₀ alkoxy group, a substituted or unsubstituted C₁-C₂₀ alkenyl group, a substituted or unsubstituted C₆-C₃₀ aryl group, a substituted or unsubstituted C₆-C₃₀ arylalkyl group, a substituted or unsubstituted C₆-C₃₀ aryloxy group, a substituted or unsubstituted C₂-C₃₀ heteroaryl group, a substituted or unsubstituted C₂-C₃₀ heteroarylalkyl group, a substituted or unsubstituted C₂-C₃₀ heteroaryloxy group, a substituted or unsubstituted C₅-C₂₀ cycloalkyl group, and a substituted or unsubstituted C₂-C₂₀ heterocycloalkyl group.

The principles of the present invention will be described with reference to FIGS. 1A and 1B, which illustrate the interaction between dimethyl oxalate as an example of an oxalate compound of formula (1) above and lithium.

Referring to FIG. 1A, in an oxalate compound of formula (1) above, two carbon-oxygen double (C=O) bonds are present in the middle portion. Accordingly,

when positive lithium ions are present around the oxalate compound of formula (1) above, localized negative charges appear between the two carbon-oxygen double bonds. Lithium ions are attracted close to the localized negative charges in the oxalate compound to form a chelate compound, as illustrated in FIG. 1B. As a result, negative sulfide ions produced by the reduction of sulfur are unlikely to bond to lithium ions, remain stable in the electrolytic solution. As a result, a reversible capacity of sulfur is increased. If lithium ions exist unbound, the free lithium ions form lithium sulfide with negative sulfide ions that is no longer available for electrochemical reactions in charging batteries.

The chelate compound illustrated in FIG. 1B migrates toward the lithium anode during charging and stays in the SEI formed on the surface of the lithium anode. The lithium ions of the chelate compound increase the conductivity of lithium ions in the SEL and stabilize the SEI.

Specified examples of an oxalate compound of formula (1) above include diethyl oxalate, dimethyl oxalate, dipropyl oxalate, dibutyl oxalate, bis-(4-methylbenzyl) oxalate, etc.

The amount of the oxalate compound of formula (1) above is in a range of 0.001-10 parts by weight, preferably 0.05-1 parts by weight, with respect to 100 parts by weight of an organic solvent of the organic electrolytic solution. If the amount of the oxalate compound of formula (1) above is less than 0.001 parts by weight, the effect of the additive is trivial. If the amount of the oxalate compound of formula (1) above exceeds 10 parts by weight, the lifespan properties of batteries degrade.

An organic solvent of an organic electrolytic solution according to the present invention is at least one selected from polyglymes of formula (2) below, dioxolanes, carbonates, 2-fluorobenzene, 3-fluorobenzene, dimethoxyethane (DME), diethoxyethane, and sulfolane (SUL).



where m is an integer from 1 to 10, and R^3 and R^4 are independently substituted or unsubstituted C_1 - C_{20} alkyl groups.

Specified examples of polyglymes of formula (2) above include diglyme (DGM, also called "diethyleneglycol dimethylether"), diethyleneglycol diethylether, triglyme

(TGM, also called "triethyleneglycol dimethylether"), triethyleneglycol diethylether, etc.

Examples of dioxolanes include 1,3-dioxolane (DOX), 4,5-diethyl-dioxolane, 4,5-dimethyl-dioxolane, 4-methyl-1,3-dioxolane, and 4-ethyl-1,3-dioxolane.

5 Examples of carbonates for the organic solvent include methylene carbonate, ethylene carbonate, diethyl carbonate, dimethyl carbonate, γ -butyrolactone, propylene carbonate, methyl ethyl carbonate, vinylene carbonate, etc.

The organic electrolytic solution according to the present invention is useful for any common lithium battery, including primary and secondary batteries,
10 especially having a lithium metal anode.

For application to lithium sulfur batteries having a cathode, which may be formed using a sulfur or a sulfur-containing compound, an organic electrolytic solution containing an oxalate compound of formula (1) above according to the present invention is prepared based on an organic solvent containing a polyglyme, a
15 dioxolane, and further at least one selected from the group consisting of sulfolane, dimethoxyethane, and diethoxyethane.

This organic solvent contains a polyglyme in an amount of 30-60% by volume, a dioxolane in an amount of 20-60% by weight, and a balance of an organic solvent selected from sulfolane, dimethoxyethane, and diethoxyethane, based on the total
20 volume of the organic solvent. It is preferable that the organic solvent contains a polyglyme and a dioxolane in a ratio of 1:9-9:1 by volume. If the amount of the dioxolane exceeds this range, discharging capacity and charging/discharging cycle life degrade.

For application of lithium batteries having a lithium composite oxide cathode,
25 an organic electrolytic solution according to the present invention is prepared based on an organic solvent containing at least one of carbonates, 2-fluorobenzene, 3-fluorobenzene, 4-fluorobenzene, dimethoxyethane, diethoxyethane, and sulfolane.

Any lithium salt which is commonly used for lithium batteries is available for an organic electrolytic solution according to the present invention. Examples of a
30 lithium salt include lithium perchlorate (LiClO_4), lithium tetrafluoroborate (LiBF_4), lithium hexafluorophosphate (LiPF_6), lithium trifluoromethanesulfonate (LiCF_3SO_3), and lithium bistrifluoromethansulfonylamide ($\text{LiN}(\text{CF}_3\text{SO}_2)_2$). It is preferable that the concentration of lithium salt is in a range of 0.5-2.0M. If the concentration of lithium salt is less than 0.5M, ionic conductivity is low. If the concentration of lithium salt

exceeds 2.0M, undesirably the lithium salt is more likely to decompose.

Examples of an unsubstituted C₁-C₂₀ alkyl group as a substituent for R₁ and R₂ in formula (1) above include a methyl group, an ethyl group, a propyl group, an isobutyl group, a sec-butyl group, a pentyl group, an iso-amyl group, a hexyl group, etc., wherein at least one hydrogen atom of the alkyl group may be substituted with halogen atom, a hydroxy group, a nitro group, a cyano group, an amino group, an amidino group, hydrazine, hydrazone, a carboxy group, a sulfonic acid group, a phosphoric acid group, a C₁-C₂₀ alkyl group, a C₂-C₂₀ alkenyl group, a C₂-C₂₀ alkynyl group, a C₁-C₂₀ heteroalkyl group, a C₆-C₂₀ aryl group, a C₆-C₂₀ arylalkyl group, a C₆-C₂₀ heteroaryl group, or a C₆-C₂₀ heteroarylalkyl group.

Examples of an unsubstituted C₁-C₂₀ alkoxy group as a substituent for R₁ and R₂ in formula (1) above include a methoxy group, an ethoxy group, a propoxy group, an isobutoxy group, a sec-butoxy group, a pentyloxy group, an iso-amyoxy group, a hexyloxy group, etc, wherein at least one hydrogen atom of the alkoxy group can be substituted with any substituent described above as being suitable for the C₁-C₂₀ alkyl group.

The aryl group as a substituent for R₁ and R₂ in formula (1) above means a C₆-C₃₀ carbocyclic aromatic system containing at least one ring wherein such rings may be attached together in a pendent manner or may be fused. The term "aryl" embraces aromatic radicals, such as phenyl, naphthyl, tetrahydronaphthyl, etc. The aryl group may have a substituent such as haloalkyl, nitro, cyano, alkoxy, and lower alkylamino. At least one hydrogen atom of the aryl group can be substituted with any substituent described above as being suitable for the C₁-C₂₀ alkyl group.

The arylalkyl group as a substituent for R₁ and R₂ in formula (1) above means the above-defined aryl group having lower alkyl substituents, for example, methyl, ethyl, propyl, etc. for some hydrogen atoms. Examples of an arylalkyl group include benzyl, phenylethyl, etc. At least one hydrogen atom of the arylalkyl group can be substituted with any substituent described above as being suitable for the C₁-C₂₀ alkyl group.

Examples of an aryloxy group as a substituent for R₁ and R₂ in formula (1) above include a phenyloxy group, a naphthyloxy group, a tetrahydronaphthyloxy group, etc. At least one hydrogen atom of the aryloxy group can be substituted with any substituent described above as being suitable for the C₁-C₂₀ alkyl group.

The heteroaryl group as a substituent for R_1 and R_2 in formula (1) above means a C_2 - C_{30} monocyclic system containing one, two, or three hetero atoms selected from the group consisting of N, O, P, and S and having at least one ring wherein such rings may be attached together in a pendent manner or may be fused. At least one hydrogen atom of the heteroaryl group can be substituted with any substituent described above as being suitable for the C_1 - C_{20} alkyl group.

The heteroarylalkyl group as a substituent for R_1 and R_2 in formula (1) above means the above-defined heteroaryl group having lower alkyl substitute groups for some hydrogen atoms, wherein at least one hydrogen atom of the heteroarylalkyl group can be substituted with any substituent described above as being suitable for the C_1 - C_{20} alkyl group.

The cycloalkyl group as a substituent for R_1 and R_2 in formula (1) above means a C_4 - C_{30} monovalent monocyclic system, wherein at least one hydrogen atom of the cycloalkyl group can be substituted with any substituent described above as being suitable for the C_1 - C_{20} alkyl group.

The heterocycloalkyl group as a substituent for R_1 and R_2 in formula (1) above means a C_1 - C_{30} monovalent monocyclic system containing one, two, or three hetero atoms selected from the group consisting of N, O, P, and S and having lower alkyl groups for some hydrogen atoms, wherein at least one hydrogen atom of the heterocycloalkyl group can be substituted with any substituent described above as being suitable for the C_1 - C_{20} alkyl group.

Hereinafter, a method of manufacturing a lithium secondary battery according to an embodiment of the present invention will be described.

First, a cathode and an anode are manufactured using a common method applied to produce lithium batteries. A lithium metal composite oxide, such as $LiCoO_2$ and $LiMn_2O_3$, is used as a cathode active material. A lithium metal electrode or a carbonaceous or graphite electrode is used for the anode.

Next, a separator is interposed between the cathode and the anode and subjected to rolling or the stacking of more electrodes and separators, to form an electrode assembly. The resulting electrode assembly is sealed in a battery case.

Next, an organic electrolytic solution according to the present invention is injected into the battery case containing the electrode assembly, so that a complete lithium secondary battery is obtained.

The organic electrolytic solution according to the present invention can be applied to lithium polymer secondary batteries, which use a polymer electrolyte, as well as such a lithium ion secondary battery described above. The organic electrolytic solution according to the present invention can be applied to lithium sulfur batteries.

Hereinafter, a method of manufacturing a lithium sulfur battery according to an embodiment of the present invention will be described. A method of manufacturing a lithium sulfur battery is similar to the above-described method of manufacturing a lithium secondary battery, with the exception of the material for the cathode. In a certain case, a protective layer may be further interposed between the anode and the separator so as to suppress the reaction of lithium with the electrolytic solution.

At least one selected from the group consisting of simple substance sulfur, kasolite containing Li_2S_n where $n \geq 1$, organo-sulfur, and $(\text{C}_2\text{S}_x)_y$ where x ranges from 2.5 to 20 and $y \geq 2$ is used as a material for the cathode. It is preferable that the anode is formed as a lithium metal electrode, a lithium-metal alloy electrode made of, for example, lithium-aluminum, lithium-magnesium or lithium-silicon, or a lithium-inert sulfur composite electrode.

The present invention will be described in greater detail with reference to the following examples. The following examples are for illustrative purposes and are not intended to limit the scope of the invention.

To measure the charging/discharging efficiency of a lithium battery containing an organic electrolytic solution according to the present invention, 2016 type half coin batteries were manufactured in Examples 1 through 13, and full lithium sulfur batteries were manufactured in Examples 14 through 19.

Example 1

An electrode assembly including a cathode, an anode, and a polyethylene separator (Ashai Co.) between the cathode and the anode was manufactured, wherein lithium metal electrodes were used for the cathode and the anode.

The electrode assembly was sealed in a battery case, and an organic electrolytic solution according to the present invention was injected to provide a complete lithium battery.

The organic electrolytic solution contained 1M LiCF_3SO_3 as a lithium salt, a

mixture of 1,3-dioxane (DOX), diglyme (DGM), dimethoxyethane (DME), and sulfolane (SUL) in a ratio of 50:20:20:10 by volume as an organic solvent, and 0.05 parts by weight of dimethyl oxalate with respect to 100 parts by weight of the organic solvent.

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Examples 2 through 4

Lithium batteries were manufactured in the same manner as in Example 1, except that the amount of dimethyl oxalate was varied to 0.1 parts by weight, 0.25 parts by weight, and 0.5 parts by weight for the respective lithium batteries.

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Example 5

A lithium battery was manufactured in the same manner as in Example 2, except that 0.1 parts by weight of diethyl oxalate was used instead of 0.1 parts by weight of dimethyl oxalate.

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Examples 6 and 7

Lithium batteries were manufactured in the same manner as in Example 5, except that the amount of diethyl oxalate was varied to 0.25 parts by weight and 0.5 parts by weight for the respective lithium batteries.

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Example 8

A lithium battery was manufactured in the same manner as in Example 2, except that 0.1 parts by weight of dibutyl oxalate was used instead of 0.1 parts by weight of dimethyl oxalate.

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Examples 9 and 10

Lithium batteries were manufactured in the same manner as in Example 8, except that the amount of dibutyl oxalate was varied to 0.25 parts by weight and 0.5 parts by weight for the respective lithium batteries.

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Example 11

A lithium battery was manufactured in the same manner as in Example 2, except that 0.1 parts by weight of bis(4-methylbenzyl) oxalate was used instead of 0.1 parts by weight of dimethyl oxalate.

Examples 12 and 13

Lithium batteries were manufactured in the same manner as in Example 11, except that the amount of bis(4-methylbenzyl) oxalate was varied to 0.25 parts by weight and 0.5 parts by weight for the respective lithium batteries.

Example 14

A cathode active material composition was prepared by mixing 70 parts by weight of sulfur, 10 parts by weight of Ketjen black as a carbon source, and 20 parts by weight of polyethyleneoxide in acetonitrile.

A carbon-coated aluminum (Al) substrate was coated with the cathode active material composition and dried to form a cathode. A lithium metal electrode was used as an anode.

An electrode assembly including the cathode, a polyethylene separator, and the anode stacked upon one another was manufactured and sealed in a battery case. An organic electrolytic solution was injected into the battery case to provide a complete lithium sulfur battery.

The organic electrolytic solution contained 1M LiCF_3SO_3 as a lithium salt, a mixture of DOX, DGM, DME, and SUL in a ratio of 50:20:20:10 by volume as an organic solvent, and 0.5 part by weight of dimethyl oxalate with respect to 100 parts by weight of the organic solvent.

Example 15

A lithium sulfur battery was manufactured in the same manner as in Example 14, except that diethyl oxalate instead of dimethyl oxalate was used for the organic electrolytic solution.

Example 16

A lithium sulfur battery was manufactured in the same manner as in Example 14, except that dibutyl oxalate instead of dimethyl oxalate was used for the organic electrolytic solution.

Example 17

A lithium sulfur battery was manufactured in the same manner as in Example

14, except that bis(4-methylbenzyl) oxalate instead of dimethyl oxalate was used for the organic electrolytic solution.

Example 18

5 A lithium sulfur battery was manufactured in the same manner as in Example 14, except that the amount of dimethyl oxalate was varied to 0.25 parts by weight.

Example 19

10 A lithium sulfur battery was manufactured in the same manner as in Example 14, except that the amount of dimethyl oxalate was varied to 1 part by weight.

Comparative Example 1

15 A lithium battery was manufactured in the same manner as in Example 1, except that only the mixture of DOX, DGM, DME, and SUL in a ratio of 50:20:20:10 by volume was used for the organic solvent, without the addition of dimethyl oxalate.

Comparative Example 2

20 A lithium battery was manufactured in the same manner as in Comparative Example 1, except that a mixture of DGM, DME, and DOX in a ratio of 40:40:20 by volume was used for the organic solvent.

Comparative Example 3

25 A lithium sulfur battery was manufactured in the same manner as in Example 14, except that only the mixture of DOX, DGM, DME, and SUL in a ratio of 50:20:20:10 by volume was used for the organic solvent, without the addition of dimethyl oxalate.

Comparative Example 4

30 A lithium sulfur battery was manufactured in the same manner as in Comparative Example 3, except that a mixture of DGM, DME, and DOX in a ratio of 40:40:20 by volume was used for the organic solvent.

Change in charging/discharging efficiency with respect to dimethyl oxalate content was measured using the lithium batteries manufactured in Examples 1 through 4 and Comparative Example 1. The results are shown in FIG. 2. As is

apparent from FIG. 2, the charging/discharging efficiency is improved for the lithium batteries of Examples 1 through 4, compared to the lithium battery of Comparative Example 1. The charging/discharging efficiency is greatest at about 0.25 parts by weight of dimethyl oxalate (Example 3).

5 Change in charging/discharging efficiency with respect to diethyl oxalate content was measured using the lithium batteries manufactured in Examples 5 through 7 and compared with Comparative Example 1. The results are shown in FIG. 3. As is apparent from FIG. 3, the charging/discharging efficiency is improved for the lithium batteries of Examples 5 through 7, compared to the lithium battery of
10 Comparative Example 1. The charging/discharging efficiency is greatest at about 0.25 parts by weight of diethyl oxalate (Example 6).

 Change in charging/discharging efficiency with respect to dibutyl oxalate content was measured using the lithium batteries manufactured in Examples 8 through 10 and compared with Comparative Example 1. The results are shown in
15 FIG. 4. As is apparent from FIG. 4, the charging/discharging efficiency is improved for the lithium batteries of Examples 8 through 10, compared to the lithium battery of Comparative Example 1. The charging/discharging efficiency is greatest at about 0.1 parts by weight of dibutyl oxalate (Example 8).

 Change in charging/discharging efficiency with respect to bis(4-methylbenzyl)
20 oxalate content was measured using the lithium batteries manufactured in Examples 11 through 13 and compared with Comparative Example 1. The results are shown in FIG. 5. As is apparent from FIG. 5, the charging/discharging efficiency is improved for the lithium batteries of Examples 11 through 13, compared to the lithium battery of Comparative Example 1. The charging/discharging efficiency is
25 greater at about 0.25 parts by weight (Example 12) and about 0.5 parts by weight (Example 13) of bis(4-methylbenzyl) oxalate.

 Change in discharging capacity with respect to the number of charging/discharging cycles was measured using the lithium sulfur batteries manufactured in Examples 14, 18 and 19 with varying amounts of dimethyl oxalate.
30 The results are shown in FIG. 6. As is apparent from FIG. 6, the discharging capacity is excellent for all of the lithium sulfur batteries of Examples 14, 18, and 19 at similar levels.

 Charging/discharging efficiency was measured using the lithium sulfur batteries manufactured in Examples 14 through 17 and Comparative Examples 1

and 2. The results are shown in FIG. 7. In FIG. 7, "A" denotes the case using the electrolytic solution of 1M LiCF_3SO_3 in the mixture of DOX, DGM, DME, and SUL in a ratio of 5:2:2:1 (Comparative Example 1), "B" denotes the case using the electrolytic solution of 1M LiCF_3SO_3 in the mixture of DGM, DME, and DOX in a ratio of 4:2:2 (Comparative Example 2), "C" denotes the case using the electrolytic solution of 1M LiCF_3SO_3 and dimethyl oxalate in the mixture of DOX, DGM, DME, and SUL in a ratio of 5:2:2:1 (Example 14), "D" denotes the case using the electrolytic solution of 1M LiCF_3SO_3 and diethyl oxide in the mixture of DOX, DGM, DME, and SUL in a ratio of 5:2:2:1 (Example 15), "E" denotes the case using the electrolytic solution of 1M LiCF_3SO_3 and dibutyl oxalate in the mixture of DOX, DGM, DME, and SUL in a ratio of 5:2:2:1 (Example 16), and "F" denotes the case using the electrolytic solution of 1M LiCF_3SO_3 and bis(4-methylbenzyl) oxalate in the mixture of DOX, DGM, DME, and SUL in a ratio of 5:2:2:1 (Example 17).

As is apparent from FIG. 7, the charging/discharging efficiency is improved for the lithium sulfur batteries of Examples 14 through 17, compared to Comparative Examples 1 and 2.

Change in discharging capacity with respect to the number of charging/discharging cycles was measured using the lithium sulfur batteries manufactured in Example 14 and Comparative Example 3. The results are shown in FIG. 8. As is apparent from FIG. 8, the discharging capacity is improved for the lithium sulfur battery of Example 14, compared to Comparative Example 3.

An organic electrolytic solution according to the present invention contains an oxalate compound of formula (1) above and thus stabilizes lithium metal and improves the conductivity of lithium ions. The organic electrolytic solution according to the present invention improves charging/discharging efficiency when used in lithium batteries having a lithium metal anode.

Especially when the organic electrolytic solution is used in lithium sulfur batteries, the oxalate compound forms a chelate with lithium ions and improves the ionic conductivity and the charging/discharging efficiency of the battery. In addition, due to the chelation of the lithium ions, negative sulfur ions remain free without interaction with lithium ions, are highly likely to dissolve in the electrolytic solution. As a result, a reversible capacity of sulfur is increased.

While the present invention has been particularly shown and described with reference to exemplary embodiments thereof, it will be understood by those of

ordinary skill in the art that various changes in form and details may be made therein without departing from the spirit and scope of the present invention as defined by the following claims.